California Medical Device Recall Information

Recall Name
GE Healthcare Recalls Aestiva/5 7900 Ventilator Due to a Potential Overdose

Recall Date	Product Description	Recalling Firm	Recall Reason
4/10/12	Aestiva/5 7900 Ventilator	**GE Healthcare, LLC.** Wauwatosa, WI	*Potential for two vaporizers to deliver each agent at the same time*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Aestiva/5 7900 Ventilator Suspect S/Ns Recalled: • AMRP01031 • AMRP00966 • AMRP01030 • AMRP00968 • AMRP00967 • AMRP01033 • AMRP00970 • AMRP00969	**CA**, nationwide	Manufactured on July 2, 2010

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm311325.htm